

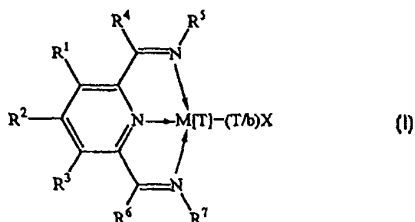
PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : C08F 10/00, 4/70	A1	(11) International Publication Number: WO 00/24788 (43) International Publication Date: 4 May 2000 (04.05.00)
<p>(21) International Application Number: PCT/GB99/03502</p> <p>(22) International Filing Date: 21 October 1999 (21.10.99)</p> <p>(30) Priority Data: 98430023.6 26 October 1998 (26.10.98) EP</p> <p>(71) Applicant (for all designated States except US): BP CHEMICALS LIMITED [GB/GB]; Britannic House, 1 Finsbury Circus, London EC2M 7BA (GB).</p> <p>(71) Applicant (for FR only): BP CHEMICALS S.N.C. [FR/FR]; Tour Neptune, La Defense, 20, place de Seine, F-92400 Courbevoie (FR).</p> <p>(72) Inventors; and (75) Inventors/Applicants (for US only): BERARDI, Alain [FR/FR]; 12, rue Braille, F-13005 Marseille (FR). SPEAKMAN, John, Gabriel [GB/FR]; 25, rue Bel Air, F-13500 Martigues (FR).</p> <p>(74) Agent: SMITH, Julian, Philip, Howard; BP International Limited, Patents & Agreements, Chertsey Road, Sunbury on Thames, Middlesex TW16 7LN (GB).</p>	<p>(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p>Published With international search report.</p>	

(54) Title: POLYMERISATION CATALYSTS



(57) Abstract

The present invention relates to a process for the polymerisation and copolymerisation of 1-olefins and to a catalyst to be used in said process wherein the catalyst comprises (1) a complex having formula (I) wherein M is Fe[II], Fe[III], Co[I], Co[II], Co[III], Mn[I], Mn[II], Mn[III], Mn[IV], Ru[II], Ru[III] or Ru[IV]; X represents an atom or group covalently or ionically bonded to the metal M; T is the oxidation state of the metal; b is the valency of the atom or group X; and R¹ to R⁷ are each independently selected from hydrogen, halogen, hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl, substituted heterohydrocarbyl or SiR'₃ where each R' is independently selected from hydrogen, halogen, hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl, substituted heterohydrocarbyl; and when any two or more of R¹ to R⁷ are hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl, said two or more can be linked to form one or more cyclic substituents; (2) an activator which is an alkylalumoxane; and (3) a support material, wherein the atomic ratio of aluminium in the alkylalumoxane (2) to transition metal M in (1) is from 6:1 to 25:1.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

POLYMERISATION CATALYSTS

The present invention relates to transition metal compounds and to polymerisation catalyst systems employing them.

The use of certain transition metal compounds to polymerise 1-olefins, for example, ethylene, is well established in the prior art. The use of Ziegler-Natta catalysts, for example, those catalysts produced by activating titanium halides with organometallic compounds such as triethylaluminium, is fundamental to many commercial processes for manufacturing polyolefins. Over the last twenty or thirty years, advances in the technology have led to the development of Ziegler-Natta catalysts which have such high activities that that olefin polymers and copolymers containing very low concentrations of residual catalyst can be produced directly in commercial polymerisation processes. The quantities of residual catalyst remaining in the produced polymer are so small as to render unnecessary their separation and removal for most commercial applications. Such processes can be operated by polymerising the monomers in the gas phase, or in solution or in suspension in a liquid hydrocarbon diluent. Polymerisation of the monomers can be carried out in the gas phase (the "gas phase process"), for example by fluidising under polymerisation conditions a bed comprising the target polyolefin powder and particles of the desired catalyst using a fluidising gas stream comprising the gaseous monomer. In the so-called "solution process" the (co)polymerisation is conducted by introducing the monomer into a solution or suspension of the catalyst in a liquid hydrocarbon diluent under conditions of temperature and pressure such that the produced polyolefin forms as a solution in the hydrocarbon diluent. In the "slurry process" the temperature, pressure and choice of diluent are such that the produced polymer forms as a suspension in the liquid hydrocarbon diluent. These processes are generally operated at relatively low pressures (for example 10-50 bar) and low temperature (for example 50 to 150°C).

Commodity polyethylenes are commercially produced in a variety of different types and grades. Homopolymerisation of ethylene with transition metal based catalysts leads to the production of so-called "high density" grades of polyethylene. These polymers have relatively high stiffness and are useful for making articles where inherent rigidity is required. Copolymerisation of ethylene with higher 1-olefins (eg butene, hexene or octene) is employed commercially to provide a wide variety of copolymers differing in density and in other important physical properties. Particularly important copolymers made by copolymerising ethylene with higher 1-olefins using transition metal based catalysts are the copolymers having a density in the range of 0.91 to 0.93. These copolymers which are generally referred to in the art as "linear low density polyethylene" are in many respects similar to the so called "low density" polyethylene produced by the high pressure free radical catalysed polymerisation of ethylene. Such polymers and copolymers are used extensively in the manufacture of flexible blown film.

An important feature of the microstructure of the copolymers of ethylene and higher 1-olefins is the manner in which polymerised comonomer units are distributed along the "backbone" chain of polymerised ethylene units. The conventional Ziegler-Natta catalysts have tended to produce copolymers wherein the polymerised comonomer units are clumped together along the chain. To achieve especially desirable film properties from such copolymers the comonomer units in each copolymer molecule are preferably not clumped together, but are well spaced along the length of each linear polyethylene chain. In recent years the use of certain metallocene catalysts (for example biscyclopentadienylzirconium dichloride activated with alumoxane) has provided catalysts with potentially high activity and capable of providing an improved distribution of the comonomer units. However, metallocene catalysts of this type suffer from a number of disadvantages, for example, high sensitivity to impurities when used with commercially available monomers, diluents and process gas streams, the need to use large quantities of expensive alumoxanes to achieve high activity, and difficulties in putting the catalyst on to a suitable support.

WO98/27124 discloses that ethylene may be polymerised by contacting it with certain iron or cobalt complexes of selected 2,6-pyridinecarboxaldehydebis(imines) and 2,6-diacetylpyridinebis(imines). These complexes are disclosed as being suitable for preparing homopolymers of ethylene. It is disclosed that in polymerisation processes, the complexes may be used in association with a neutral Lewis acid such as

methyaluminoxane (MAO). Ratios of aluminium in the MAO to Fe or Co in the complex exemplified are in the range from 31:1 to 2485:1.

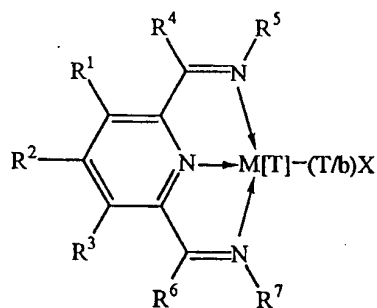
We have developed novel catalysts utilising complexes similar to the above which are disclosed in our copending application WO GB98/2638. This discloses

5 polymerisation of ethylene using a catalyst which comprises a transition metal salt of a 2,6-diacylpyridinebis(imine) supported on silica, with an MAO cocatalyst. Ratios of aluminium in the MAO to transition metal in the complex are stated to range from 0.1-20000:1, preferably 1-2000:1, and typically at least 500:1. Ratios in the examples range from 31:1 upwards.

10 We have now surprisingly discovered that the activity of catalysts comprising the above compounds and MAO in polymerisation of 1-olefins can be maintained or even improved by reducing the ratio of aluminium in the MAO to transition metal.

Accordingly in a first aspect the invention provides a catalyst for the polymerisation of olefins comprising

15 (1) a complex having the formula (I)



Formula (I)

wherein M is Fe[II], Fe[III], Co[I], Co[II], Co[III], Mn[I], Mn[II], Mn[III], Mn[IV],
 20 Ru[II], Ru[III] or Ru[IV]; X represents an atom or group covalently or ionically bonded to the metal M; T is the oxidation state of the metal; b is the valency of the atom or group X; and R¹ to R⁷ are each independently selected from hydrogen, halogen, hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl, substituted heterohydrocarbyl or SiR'₃, where each R' is independently selected from hydrogen, halogen, hydrocarbyl,
 25 substituted hydrocarbyl, heterohydrocarbyl, substituted heterohydrocarbyl; and when any two or more of R¹ to R⁷ are hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl, said two or more can be linked to form one or more cyclic

substituents;

(2) an activator which is an alkylalumoxane; and

(3) a support material,

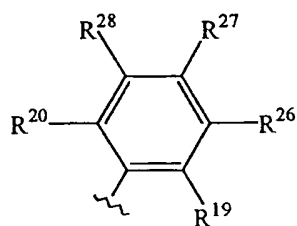
wherein the atomic ratio of aluminium in the alkylalumoxane (2) to transition metal M in

5 (1) is from 6:1 to 25:1.

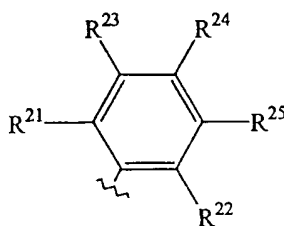
As alkylalumoxanes are added to catalysts in order to enhance their activity, the invention is particularly surprising. Alkylalumoxanes are relatively expensive and also undesirable from a safety point of view, so the ability to use smaller quantities whilst maintaining or enhancing polymerisation activity is especially advantageous. It is
10 preferred that the atomic ratio of aluminium to transition metal M is from 8:1 to 22:1, and more preferably from 12:1 to 18:1.

The activator (2) is preferably a (C₁-C₄) alkylalumoxane, the alkyl group generally being methyl, ethyl, propyl or isobutyl. Preferred is methylalumoxane (also known as methylaluminumoxane or MAO) or modified methylalumoxane (MMAO), which
15 additionally contains isobutylalumoxane. The term "alkylalumoxane" as used in this specification includes alkylalumoxanes available commercially which may contain a proportion, typically about 10wt%, but optionally up to 50wt%, of the corresponding trialkylaluminium; for instance, commercial MAO usually contains approximately 10% trimethylaluminium (TMA), whilst commercial MMAO contains both TMA and
20 triisobutylaluminium (TIBA). Quantities of alkylalumoxane quoted herein include such trialkylaluminium impurities, and accordingly the amount of aluminium used to calculate the above ratio in this invention is based on the amount present in all 'components' of the alkylalumoxane.

In a preferred complex of Formula (I), R⁵ is represented by the group "P" and R⁷
25 is represented by the group "Q" as follows:



Group P



Group Q

wherein R^{19} to R^{28} are independently selected from hydrogen, halogen, hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl; when any two or more of R^1 to R^4 , R^6 and R^{19} to R^{28} are hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl, said two or more can be linked to
 5 form one or more cyclic substituents.

The ring systems P and Q are preferably independently 2,6-hydrocarbylphenyl or fused-ring polyaromatic, for example, 1-naphthyl, 2-naphthyl, 1-phenanthrenyl and 8-quinoliny.

Preferably at least one of R^{19} , R^{20} , R^{21} and R^{22} is hydrocarbyl, substituted
 10 hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl. More preferably at least one of R^{19} and R^{20} , and at least one of R^{21} and R^{22} , is hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl. Most preferably R^{19} , R^{20} , R^{21} and R^{22} are all independently selected from hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl. R^{19} , R^{20} , R^{21} and R^{22} are preferably
 15 independently selected from methyl, ethyl, n-propyl, iso-propyl, n-butyl, sec-butyl, tert.-butyl, n-pentyl, neopentyl, n-hexyl, 4-methylpentyl, n-octyl, phenyl and benzyl.

R^1 , R^2 , R^3 , R^4 , R^6 , R^{19} , R^{20} , R^{21} , R^{22} , R^{23} , R^{25} , R^{26} and R^{28} are preferably independently selected from hydrogen and C_1 to C_8 hydrocarbyl, for example, methyl, ethyl, n-propyl, n-butyl, t-butyl, n-hexyl, n-octyl, phenyl and benzyl.

20 R^{24} and R^{27} are preferably each independently selected from ethyl, n-propyl, iso-propyl, n-butyl, sec-butyl, tert.-butyl, n-pentyl, neopentyl, n-hexyl, 4-methylpentyl, n-octyl, phenyl and benzyl, or alternatively are each independently fluorine, chlorine, bromine or iodine, optionally both fluorine.

In an alternative embodiment R^5 is a group having the formula $-NR^{29}R^{30}$ and R^7
 25 is a group having the formula $-NR^{31}R^{32}$, wherein R^{29} to R^{32} are independently selected from hydrogen, halogen, hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl; when any two or more of R^1 to R^4 , R^6 and R^{29} to R^{32} are hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl, said two or more can be linked to form one or more cyclic
 30 substituents.

Each of the nitrogen atoms is coordinated to the metal by a "dative" bond, ie a bond formed by donation of a lone pair of electrons from the nitrogen atom. The remaining bonds on each of these atoms are covalent bonds formed by electron sharing

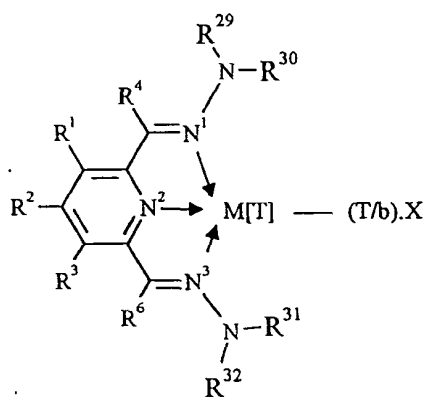
between the atoms and the organic ligand as shown in the defined formula for the metal complex illustrated above.

Preferably in Formula B above M is Fe[II], Fe[III], Ru[II], Mn[II], Co[II], Ru[III] or Ru[IV].

5 Yet further compounds suitable for the catalyst systems of the present invention are those comprising the skeletal unit depicted in Formula T:

10

15



Formula T

20

wherein M is Fe[II], Fe[III], Co[I], Co[II], Co[III], Mn[I], Mn[II], Mn[III], Mn[IV], Ru[II], Ru[III] or Ru[IV]; X represents an atom or group covalently or ionically bonded to the transition metal M; T is the oxidation state of the transition metal M and b is the valency of the atom or group X; R¹ to R⁴, R⁶ and R²⁹ to R³² are independently selected from hydrogen, halogen, hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl; when any two or more of R¹ to R⁴, R⁶ and R²⁹ to R³² are hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl, said two or more can be linked to form one or more cyclic substituents.

30

The atom or group represented by X in the compounds of Formula (I) and T can be, for example, selected from halide, sulphate, nitrate, thiolate, thiocarboxylate, BF₄⁻, PF₆⁻, hydride, hydrocarbyloxy, carboxylate, hydrocarbyl, substituted hydrocarbyl and

heterohydrocarbyl, or β -diketonates. Examples of such atoms or groups are chloride, bromide, methyl, ethyl, propyl, butyl, octyl, decyl, phenyl, benzyl, methoxide, ethoxide, isopropoxide, tosylate, triflate, formate, acetate, phenoxide and benzoate. Preferred examples of the atom or group X in the compounds of Formula (I) are halide, for example, chloride, bromide; hydride; hydrocarbyloxy, for example, methoxide, ethoxide, isopropoxide, phenoxide; carboxylate, for example, formate, acetate, benzoate; hydrocarbyl, for example, methyl, ethyl, propyl, butyl, octyl, decyl, phenyl, benzyl; substituted hydrocarbyl; heterohydrocarbyl; tosylate; and triflate. Preferably X is selected from halide, hydride and hydrocarbyl. Chloride is particularly preferred.

In the compound of Formula (I) of the present invention, M is preferably Fe[II]. In the compounds of Formula T of the present invention, M is preferably Fe[II], Mn[II] or Co[II].

The following are examples of nitrogen-containing transition metal complexes that can be employed in the catalyst of the present invention:

- 2,6-diacetylpyridinebis(2,6-diisopropylanil)FeCl₂
- 2,6-diacetylpyridine(2,6-diisopropylanil)MnCl₂
- 2,6-diacetylpyridine(2,6-diisopropylanil)CoCl₂
- 2,6-diacetylpyridinebis(2-tert.-butylanil)FeCl₂
- 2,6-diacetylpyridinebis(2,3-dimethylanil)FeCl₂
- 2,6-diacetylpyridinebis(2-methylanil)FeCl₂
- 2,6-diacetylpyridinebis(2,4-dimethylanil)FeCl₂
- 2,6-diacetylpyridinebis(2,6-dimethylanil)FeCl₂
- 2,6-diacetylpyridinebis(2,4,6-trimethylanil)FeCl₂
- 2,6-dialdiminepyridinebis(2,6-dimethylanil)FeCl₂
- 2,6-dialdiminepyridinebis(2,6-diethylanil)FeCl₂
- 2,6-dialdiminepyridinebis(2,6-diisopropylanil)FeCl₂
- 2,6-dialdiminepyridinebis(1-naphthyl)FeCl₂ and
- 2,6-bis(1,1-diphenylhydrazon)pyridine.FeCl₂.

A preferred complex of the present invention is 2,6-diacetylpyridinebis(2,4,6-trimethylanil)FeCl₂.

The polymerisation catalyst system of the invention may additionally comprise (3) a neutral Lewis base. Neutral Lewis bases are well known in the art of Ziegler-Natta catalyst polymerisation technology. Examples of classes of neutral Lewis bases suitably

employed in the present invention are unsaturated hydrocarbons, for example, alkenes (other than 1-olefins) or alkynes, primary, secondary and tertiary amines, amides, phosphoramides, phosphines, phosphites, ethers, thioethers, nitriles, carbonyl compounds, for example, esters, ketones, aldehydes, carbon monoxide and carbon dioxide, sulphoxides, sulphones and boroxines. Although 1-olefins are capable of acting as neutral Lewis bases, for the purposes of the present invention they are regarded as monomer or comonomer 1-olefins and not as neutral Lewis bases *per se*. However, alkenes which are internal olefins, for example, 2-butene and cyclohexene are regarded as neutral Lewis bases in the present invention. Preferred Lewis bases are tertiary amines and aromatic esters, for example, dimethylaniline, diethylaniline, tributylamine, ethylbenzoate and benzylbenzoate. In this particular aspect of the present invention, components (1), (2) and (3) of the catalyst system can be brought together simultaneously or in any desired order. However, if components (2) and (3) are compounds which interact together strongly, for example, form a stable compound together, it is preferred to bring together either components (1) and (2) or components (1) and (3) in an initial step before introducing the final defined component. Preferably components (1) and (3) are contacted together before component (2) is introduced. The quantities of components (1) and (2) employed in the preparation of this catalyst system are suitably as described above in relation to the catalysts of the present invention. The quantity of the neutral Lewis Base [component (3)] is preferably such as to provide a ratio of component (1):component (3) in the range 100:1 to 1:1000, most preferably in the range 1:1 to 1:20. Components (1), (2) and (3) of the catalyst system can brought together, for example, as the neat materials, as a suspension or solution of the materials in a suitable diluent or solvent (for example a liquid hydrocarbon), or, if at least one of the components is volatile, by utilising the vapour of that component. The components can be brought together at any desired temperature. Mixing the components together at room temperature is generally satisfactory. Heating to higher temperatures e.g. up to 120°C can be carried out if desired, e.g. to achieve better mixing of the components. It is preferred to carry out the bringing together of components (1), (2) and (3) in an inert atmosphere (e.g. dry nitrogen) or *in vacuo*. If it is desired to use the catalyst on a support material (see below), this can be achieved, for example, by preforming the catalyst system comprising components (1), (2) and (3) and impregnating the support material preferably with a solution thereof, or by introducing to the support material one

or more of the components simultaneously or sequentially. If desired the support material itself can have the properties of a neutral Lewis base and can be employed as, or in place of, component (3). An example of a support material having neutral Lewis base properties is poly(aminostyrene) or a copolymer of styrene and aminostyrene (ie
5 vinylaniline).

The catalysts of the present invention can if desired comprise more than one of the defined compounds. Alternatively, the catalysts of the present invention can also include one or more other types of transition metal compounds or catalysts, for example, nitrogen containing catalysts such as those described in our copending applications
10 PCT/GB98/02638 or GB 9903402.7. Examples of such other catalysts include 2,6-diacetylpyridinebis(2,4,6-trimethyl anil)FeCl₂.

The catalysts of the present invention can also include one or more other types of catalyst, such as those of the type used in conventional Ziegler-Natta catalyst systems, metallocene-based catalysts, monocyclopentadienyl- or constrained geometry based
15 catalysts, or heat activated supported chromium oxide catalysts (eg Phillips-type catalyst).

The catalysts of the present invention can be unsupported or supported on a support material, for example, silica, alumina, MgCl₂ or zirconia, or on a polymer or prepolymer, for example polyethylene, polypropylene, polystyrene, or
20 poly(aminostyrene).

If desired the catalysts can be formed in situ in the presence of the support material, or the support material can be pre-impregnated or premixed, simultaneously or sequentially, with one or more of the catalyst components. The catalysts of the present invention can if desired be supported on a heterogeneous catalyst, for example, a
25 magnesium halide supported Ziegler Natta catalyst, a Phillips type (chromium oxide) supported catalyst or a supported metallocene catalyst. Formation of the supported catalyst can be achieved for example by treating the transition metal compounds of the present invention with alumoxane in a suitable inert diluent, for example a volatile hydrocarbon, slurring a particulate support material with the product and evaporating
30 the volatile diluent. The produced supported catalyst is preferably in the form of a free-flowing powder. The quantity of support material employed can vary widely, for example from 100,000 to 1 grams per gram of metal present in the transition metal compound.

The present invention further provides a process for the polymerisation and

copolymerisation of 1-olefins, comprising contacting the monomeric olefin under polymerisation conditions with the polymerisation catalyst or catalyst system of the present invention. A preferred process comprises the steps of :

- a) preparing a prepolymer-based catalyst by contacting one or more 1-olefins with a catalyst system, and
- b) contacting the prepolymer-based catalyst with one or more 1-olefins, wherein the catalyst system is as defined above.

In the text hereinbelow, the term "catalyst" is intended to include "catalyst system" as defined previously and also "prepolymer-based catalyst" as defined above.

The polymerisation conditions can be, for example, solution phase, slurry phase, gas phase or bulk phase, with polymerisation temperatures ranging from -100°C to +300°C, and at pressures of atmospheric and above, particularly from 140 to 4100 kPa. If desired, the catalyst can be used to polymerise ethylene under high pressure/high temperature process conditions wherein the polymeric material forms as a melt in supercritical ethylene. Preferably the polymerisation is conducted under gas phase fluidised bed or stirred bed conditions.

Prior to the polymerisation, further support material impregnated with a compound of the formula MR_3 , where each R is independently C_1 - C_{12} alkyl or halo, and M is Al, Li, B, Mg or Zn, may be added to the reactor vessel; this acts as a poison scavenger. The above calculation of Al:M ratio does not include any Al impregnated in this additional support material.

Optionally, a compound of the formula AlR_3 , where each R is independently C_1 - C_{12} alkyl or halo, may be separately added to the reactor. Again, the aluminium thereby added is not included in the calculation of the above ratio. The three substituents R in the compound of the formula AlR_3 , which may be the same or different, are preferably hydrogen, methyl, ethyl, butyl or chloro. Preferred compounds AlR_3 include trimethylaluminium (TMA), triethylaluminium (TEA), tri-isobutylaluminium (TIBA), tri-n-octylaluminium, ethylaluminium dichloride and diethylaluminium chloride. Most preferred are TMA and TIBA.

Suitable monomers for use in the polymerisation process of the present invention are, for example, ethylene and C_{2-20} α -olefins, specifically propylene, 1-butene, 1-pentene, 1-hexene, 4-methylpentene-1, 1-heptene, 1-octene, 1-nonene, 1-decene, 1-undecene, 1-dodecene, 1-tridecene, 1-tetradecene, 1-pentadecene, 1-hexadecene, 1-

heptadecene, 1-octadecene, 1-nonadecene, and 1-eicosene. Other monomers include methyl methacrylate, methyl acrylate, butyl acrylate, acrylonitrile, vinyl acetate, and styrene. Preferred monomers for homopolymerisation processes are ethylene and propylene.

5 The catalysts and process of the invention can also be used for copolymerising ethylene or propylene with each other or with other 1-olefins such as 1-butene, 1-hexene, 4-methylpentene-1, and octene, or with other monomeric materials, for example, methyl methacrylate, methyl acrylate, butyl acrylate, acrylonitrile, vinyl acetate, and styrene.

Irrespective of the polymerisation or copolymerisation technique employed,
10 polymerisation or copolymerisation is typically carried out under conditions that substantially exclude oxygen, water, and other materials that act as catalyst poisons. Also, polymerisation or copolymerisation can be carried out in the presence of additives to control polymer or copolymer molecular weights.

The use of hydrogen gas as a means of controlling the average molecular weight
15 of the polymer or copolymer applies generally to the polymerisation process of the present invention. For example, hydrogen can be used to reduce the average molecular weight of polymers or copolymers prepared using gas phase, slurry phase, bulk phase or solution phase polymerisation conditions. The quantity of hydrogen gas to be employed to give the desired average molecular weight can be determined by simple "trial and
20 error" polymerisation tests.

The polymerisation process of the present invention provides polymers and copolymers, especially ethylene polymers, at remarkably high productivity (based on the amount of polymer or copolymer produced per unit weight of complex employed in the catalyst system). This means that relatively very small quantities of transition metal
25 complex are consumed in commercial processes using the process of the present invention. It also means that when the polymerisation process of the present invention is operated under polymer recovery conditions that do not employ a catalyst separation step, thus leaving the catalyst, or residues thereof, in the polymer (e.g. as occurs in most commercial slurry and gas phase polymerisation processes), the amount of transition
30 metal complex in the produced polymer can be very small.

Slurry phase polymerisation conditions or gas phase polymerisation conditions are particularly useful for the production of high or low density grades of polyethylene, and polypropylene. In these processes the polymerisation conditions can be batch,

continuous or semi-continuous. Furthermore, one or more reactors may be used, e.g. from two to five reactors in series. Different reaction conditions, such as different temperatures or hydrogen concentrations may be employed in the different reactors. In the slurry phase process and the gas phase process, the catalyst is generally metered and transferred into the polymerisation zone in the form of a particulate solid either as a dry powder (e.g. with an inert gas) or as a slurry. This solid can be, for example, a solid catalyst system formed from the one or more of complexes of the invention and an activator with or without other types of catalysts, or can be the solid catalyst alone with or without other types of catalysts. In the latter situation, the activator can be fed to the polymerisation zone, for example as a solution, separately from or together with the solid catalyst. Preferably the catalyst system or the transition metal complex component of the catalyst system employed in the slurry polymerisation and gas phase polymerisation is supported on one or more support materials. Most preferably the catalyst system is supported on the support material prior to its introduction into the polymerisation zone. Suitable support materials are, for example, silica, alumina, zirconia, talc, kieselguhr, or magnesia. Impregnation of the support material can be carried out by conventional techniques, for example, by forming a solution or suspension of the catalyst components in a suitable diluent or solvent, and slurrying the support material therewith. The support material thus impregnated with catalyst can then be separated from the diluent for example, by filtration or evaporation techniques. Once the polymer product is discharged from the reactor, any associated and absorbed hydrocarbons are substantially removed, or degassed, from the polymer by, for example, pressure let-down or gas purging using fresh or recycled steam, nitrogen or light hydrocarbons (such as ethylene). Recovered gaseous or liquid hydrocarbons may be recycled to the polymerisation zone.

In the slurry phase polymerisation process the solid particles of catalyst, or supported catalyst, are fed to a polymerisation zone either as dry powder or as a slurry in the polymerisation diluent. The polymerisation diluent is compatible with the polymer(s) and catalyst(s), and may be an alkane such as hexane, heptane, isobutane, or a mixture of hydrocarbons or paraffins. Preferably the particles are fed to a polymerisation zone as a suspension in the polymerisation diluent. The polymerisation zone can be, for example, an autoclave or similar reaction vessel, or a continuous loop reactor, e.g. of the type well-known in the manufacture of polyethylene by the Phillips Process. When the polymerisation process of the present invention is carried out under slurry conditions the

polymerisation is preferably carried out at a temperature above 0°C, most preferably above 15°C. The polymerisation temperature is preferably maintained below the temperature at which the polymer commences to soften or sinter in the presence of the polymerisation diluent. If the temperature is allowed to go above the latter temperature, fouling of the reactor can occur. Adjustment of the polymerisation within these defined temperature ranges can provide a useful means of controlling the average molecular weight of the produced polymer. A further useful means of controlling the molecular weight is to conduct the polymerisation in the presence of hydrogen gas which acts as chain transfer agent. Generally, the higher the concentration of hydrogen employed, the lower the average molecular weight of the produced polymer.

In bulk polymerisation processes, liquid monomer such as propylene is used as the polymerisation medium.

Methods for operating gas phase polymerisation processes are well known in the art. Such methods generally involve agitating (e.g. by stirring, vibrating or fluidising) a bed of catalyst, or a bed of the target polymer (i.e. polymer having the same or similar physical properties to that which it is desired to make in the polymerisation process) containing a catalyst, and feeding thereto a stream of monomer at least partially in the gaseous phase, under conditions such that at least part of the monomer polymerises in contact with the catalyst in the bed. The bed is generally cooled by the addition of cool gas (e.g. recycled gaseous monomer) and/or volatile liquid (e.g. a volatile inert hydrocarbon, or gaseous monomer which has been condensed to form a liquid). The polymer produced in, and isolated from, gas phase processes forms directly a solid in the polymerisation zone and is free from, or substantially free from liquid. As is well known to those skilled in the art, if any liquid is allowed to enter the polymerisation zone of a gas phase polymerisation process the quantity of liquid in the polymerisation zone is small in relation to the quantity of polymer present.. This is in contrast to "solution phase" processes wherein the polymer is formed dissolved in a solvent, and "slurry phase" processes wherein the polymer forms as a suspension in a liquid diluent.

The gas phase process can be operated under batch, semi-batch, or so-called "continuous" conditions. It is preferred to operate under conditions such that monomer is continuously recycled to an agitated polymerisation zone containing polymerisation catalyst, make-up monomer being provided to replace polymerised monomer, and continuously or intermittently withdrawing produced polymer from the polymerisation

zone at a rate comparable to the rate of formation of the polymer, fresh catalyst being added to the polymerisation zone to replace the catalyst withdrawn from the polymerisation zone with the produced polymer.

For typical production of impact copolymers, homopolymer formed from the first monomer in a first reactor is reacted with the second monomer in a second reactor. For manufacture of propylene/ethylene impact copolymer in a gas-phase process, propylene is polymerized in a first reactor; reactive polymer transferred to a second reactor in which ethylene or other comonomer is added. The result is an intimate mixture of a isotactic polypropylene chains with chains of a random propylene/ethylene copolymer. A random copolymer typically is produced in a single reactor in which a minor amount of a comonomer (typically ethylene) is added to polymerizing chains of propylene.

Methods for operating gas phase fluidised bed processes for making polyethylene, ethylene copolymers and polypropylene are well known in the art. The process can be operated, for example, in a vertical cylindrical reactor equipped with a perforated distribution plate to support the bed and to distribute the incoming fluidising gas stream through the bed. The fluidising gas circulating through the bed serves to remove the heat of polymerisation from the bed and to supply monomer for polymerisation in the bed. Thus the fluidising gas generally comprises the monomer(s) normally together with some inert gas (e.g. nitrogen or inert hydrocarbons such as methane, ethane, propane, butane, pentane or hexane) and optionally with hydrogen as molecular weight modifier. The hot fluidising gas emerging from the top of the bed is led optionally through a velocity reduction zone (this can be a cylindrical portion of the reactor having a wider diameter) and, if desired, a cyclone and or filters to disentrain fine solid particles from the gas stream. The hot gas is then led to a heat exchanger to remove at least part of the heat of polymerisation. Catalyst is preferably fed continuously or at regular intervals to the bed. At start up of the process, the bed comprises fluidisable polymer which is preferably similar to the target polymer. Polymer is produced continuously within the bed by the polymerisation of the monomer(s). Preferably means are provided to discharge polymer from the bed continuously or at regular intervals to maintain the fluidised bed at the desired height. The process is generally operated at relatively low pressure, for example, at 10 to 50 bars, and at temperatures for example, between 50 and 120 °C. The temperature of the bed is maintained below the sintering temperature of the fluidised polymer to avoid problems of

agglomeration.

In the gas phase fluidised bed process for polymerisation of olefins the heat evolved by the exothermic polymerisation reaction is normally removed from the polymerisation zone (i.e. the fluidised bed) by means of the fluidising gas stream as described above. The hot reactor gas emerging from the top of the bed is led through one or more heat exchangers wherein the gas is cooled. The cooled reactor gas, together with any make-up gas, is then recycled to the base of the bed. In the gas phase fluidised bed polymerisation process of the present invention it is desirable to provide additional cooling of the bed (and thereby improve the space time yield of the process) by feeding a volatile liquid to the bed under conditions such that the liquid evaporates in the bed thereby absorbing additional heat of polymerisation from the bed by the "latent heat of evaporation" effect. When the hot recycle gas from the bed enters the heat exchanger, the volatile liquid can condense out. In one embodiment of the present invention the volatile liquid is separated from the recycle gas and reintroduced separately into the bed. Thus, for example, the volatile liquid can be separated and sprayed into the bed. In another embodiment of the present invention the volatile liquid is recycled to the bed with the recycle gas. Thus the volatile liquid can be condensed from the fluidising gas stream emerging from the reactor and can be recycled to the bed with recycle gas, or can be separated from the recycle gas and then returned to the bed.

The method of condensing liquid in the recycle gas stream and returning the mixture of gas and entrained liquid to the bed is described in EP-A-0089691 and EP-A-0241947. It is preferred to reintroduce the condensed liquid into the bed separate from the recycle gas using the process described in our US Patent 5541270, the teaching of which is hereby incorporated into this specification.

When using the catalysts of the present invention under gas phase polymerisation conditions, the catalyst, or one or more of the components employed to form the catalyst can, for example, be introduced into the polymerisation reaction zone in liquid form, for example, as a solution in an inert liquid diluent. Thus, for example, the transition metal component, or the activator component, or both of these components can be dissolved or slurried in a liquid diluent and fed to the polymerisation zone. Under these circumstances it is preferred the liquid containing the component(s) is sprayed as fine droplets into the polymerisation zone. The droplet diameter is preferably within the range 1 to 1000 microns. EP-A-0593083, the teaching of which is hereby incorporated

into this specification, discloses a process for introducing a polymerisation catalyst into a gas phase polymerisation. The methods disclosed in EP-A-0593083 can be suitably employed in the polymerisation process of the present invention if desired.

Although not usually required, upon completion of polymerisation or
5 copolymerisation, or when it is desired to terminate polymerisation or copolymerisation or at least temporarily deactivate the catalyst or catalyst component of this invention, the catalyst can be contacted with water, alcohols, acetone, or other suitable catalyst deactivators a manner known to persons of skill in the art.

Homopolymerisation of ethylene with the catalysts of the invention may produce
10 so-called "high density" grades of polyethylene. These polymers have relatively high stiffness and are useful for making articles where inherent rigidity is required. Copolymerisation of ethylene with higher 1-olefins (eg butene, hexene or octene) can provide a wide variety of copolymers differing in density and in other important physical properties. Particularly important copolymers made by copolymerising ethylene with
15 higher 1-olefins with the catalysts of the invention are the copolymers having a density in the range of 0.91 to 0.93. These copolymers which are generally referred to in the art as linear low density polyethylene, are in many respects similar to the so called low density polyethylene produced by the high pressure free radical catalysed polymerisation of ethylene. Such polymers and copolymers are used extensively in the manufacture of
20 flexible blown film.

Propylene polymers produced by the process of the invention include propylene homopolymer and copolymers of propylene with less than 50 mole % ethylene or other alpha-olefin such as butene-1, pentene-1, 4-methylpentene-1, or hexene-1, or mixtures thereof. Propylene polymers also may include copolymers of propylene with minor
25 amounts of a copolymerizable monomer. Typically, most useful are normally-solid polymers of propylene containing polypropylene crystallinity, random copolymers of propylene with up to about 10 wt.% ethylene, and impact copolymers containing up to about 20 wt.% ethylene or other alpha-olefin. Polypropylene homopolymers may contain a small amount (typically below 2 wt.%) of other monomers to the extent the properties
30 of the homopolymer are not affected significantly.

Propylene polymers may be produced which are normally solid, predominantly isotactic, poly α -olefins. Levels of stereorandom by-products are sufficiently low so that useful products can be obtained without separation thereof. Typically, useful propylene

homopolymers show polypropylene crystallinity and have isotactic indices above 90 and many times above 95. Copolymers typically will have lower isotactic indices, typically above 80-85.

Depending upon polymerisation conditions known in the art, propylene polymers with melt flow rates from below 1 to above 1000 may be produced in a reactor. For many applications, polypropylenes with a MFR from 2 to 100 are typical. Some uses such as for spunbonding may use a polymer with an MFR of 500 to 2000.

Peroxide compounds may be added to ethylene or propylene polymers. For ethylene based polymers, peroxides can be used to give cross-linking in the polymer. For the preparation of high MFR propylene polymers, peroxide compounds may be added during extrusion for controlled rheology to increase the melt flow rate of polymer. Peroxide acts to break long polymer chains and has the effect of both increasing MFR and narrowing the molecular weight distribution (M_w/M_n) or polydispersity. A typical reactor polypropylene powder with an MFR of 2g/10 min. by controlled rheology treatment with peroxide in an extruder may form a polymer with an MFR of 20-40. By varying the type, amount of, and process conditions using, peroxide, the final polymer MFR may be controlled as known in the art.

Depending upon the use of the polymer product, minor amounts of additives are typically incorporated into the polymer formulation such as acid scavengers, antioxidants, stabilizers, and the like. Generally, these additives are incorporated at levels of about 25 to 2000 ppm, typically from about 50 to about 1000 ppm, and more typically 400 to 1000 ppm, based on the polymer.

In use, polymers or copolymers made according to the invention in the form of a powder are conventionally compounded into pellets. Examples of uses for polymer compositions made according to the invention include use to form fibres, extruded films, tapes, spunbonded webs, moulded or thermoformed products, and the like. The polymers may be blown into films, or may be used for making a variety of moulded or extruded articles such as pipes, and containers such as bottles or drums. Specific additive packages for each application may be selected as known in the art. Examples of supplemental additives include slip agents, anti-blocks, anti-stats, mould release agents, primary and secondary anti-oxidants, clarifiers, nucleants, uv stabilizers, and the like. Classes of additives are well known in the art and include phosphite antioxidants, hydroxylamine (such as N,N-dialkyl hydroxylamine) and amine oxide (such as dialkyl methyl amine

oxide) antioxidants, hindered amine light (uv) stabilizers, phenolic stabilizers, benzofuranone stabilizers, and the like. Various olefin polymer additives are described in U.S. patents 4,318,845, 4,325,863, 4,590,231, 4,668,721, 4,876,300, 5,175,312, 5,276,076, 5,326,802, 5,344,860, 5,596,033, and 5,625,090.

- 5 Fillers such as silica, glass fibers, talc, and the like, nucleating agents, and colourants also may be added to the polymer compositions as known by the art.

The present invention is illustrated in the following Examples.

EXAMPLES

- 10 Example 1a - Preparation of 2,6-diacetylpyridine bis (2,4,6-trimethylphenyl amine) FeCl₂ complex

4g of 2,6-diacetylpyridine, 10.32ml of 2,4,6-trimethylaniline and 0.1g of p-toluenesulphonic acid monohydrate were added to 300ml of dry toluene in a 500ml round bottomed flask. A 20ml Dean-Stark heavy fractionating column and condenser
15 were attached and the mixture heated with stirring to 160°C. The reaction appeared to have gone to completion in about 3 hours. The solution was then evacuated and 200ml of methanol added. 2,6-diacetylpyridine bis (2,4,6-trimethylphenyl amine) precipitated out and was isolated by filtration and washed with 3 aliquots of 20ml of methanol.

3.19g of iron(II) chloride were weighed into a half litre Schlenk tube and 400 ml
20 of n-butanol added. The suspension was warmed to 90°C and stirred for 3 hours. 10g of 2,6-diacetylpyridine bis (2,4,6-trimethylphenyl amine) was added as a solid at 80°C. The reaction was stirred for 1 hour then at 25°C over 3 days. The resultant blue suspension was allowed to settle and the supernatant liquid decanted off. Two washes of the precipitate with 200ml aliquots of toluene were carried out, followed by one wash
25 with 100ml of n-pentane. The residual blue solid was pumped in vacuo for 6 hours till dry. It was stored and handled in a nitrogen atmosphere.

Example 1b - Preparation of the catalyst

The equipment used for calcining the silica support was composed essentially of a
30 vertical quartz cylinder, 75cm high and 5cm in diameter, above which was placed a disengagement zone. This calcinator was equipped with a fluidisation grid made from porous quartz and positioned in the lower part of the quartz cylinder. It was also equipped with an electrical resistance heater and a supply of fluidising nitrogen.

Into the calcinator maintained at 60°C and supplied with nitrogen containing less than 2 vpm of water vapour and with a flow rate of 4.7 ml/s, were charged 30g of silica sold under the trade name of ES70X by Crosfield Catalysts (Warrington, England).

Next the calcinator was heated from 60 to 600°C at a rate of 100°C/h. The silica was
5 then maintained at 600°C for 16 hours in the fluidised state. The silica was then cooled to 25°C and stored under dry nitrogen.

All subsequent operations were carried out in a nitrogen atmosphere. 12.5g of the silica thus dried were suspended in a solution of 0.66g of methylaluminoxane (Witco, Bergkamen, Germany) in 11.1g of toluene. The suspension was agitated for 2 hours at
10 25°C. The silica thus treated was washed five times with 20 ml of toluene. The treated silica was then dried under vacuum.

2.89g of the treated silica were suspended in 20ml of toluene and 0.1932g of the above iron complex (see 1a) in 10ml of toluene was added. The suspension was agitated for 17 hours at 25°C. The catalyst thus prepared was washed five times with 20 ml of toluene,
15 and then dried under vacuum.

The catalyst thus prepared contained 0.419 % by weight of iron and 2.88 % by weight of aluminium. This corresponds to an atomic ratio Al/Fe of 14.3.

Example 1c - Ethylene polymerisation in the gas phase

20 400g of polyethylene pellets were introduced into a stainless steel reactor of capacity 2.6 litres equipped with a stirrer and maintained under a nitrogen atmosphere. The reactor was heated to 100°C, stirred at 50 rpm and 0.7g of silica previously treated with 1.5mmole/g of triethylaluminium was added as poison scavenger. Next 0.1025g of the above catalyst (see 1b) was added. Then hydrogen was introduced to pressure the
25 reactor to 0.1 MPa. Finally ethylene was fed until a total pressure of 0.8 MPa was obtained. Ethylene was fed to maintain this pressure throughout the reaction. After production corresponding to 50g polyethylene per gram of catalyst, the stirring rate was increased to 100rpm, after the production of 50g of polyethylene the stirring rate was increased further to 150 rpm and after the production of 100g of polyethylene the stirring
30 rate was increased further to 200 rpm.

Polymerisation was allowed to continue for 1 hour and 18 minutes, during which period 223g of polyethylene were produced. This corresponds to an average activity of 3372g/mmol Fe/h/bar. The reactor contents were cooled to 25°C and were then

recovered from the reactor. A sieve with 2mm diameter mesh was used to separate the polyethylene pellets from the powder formed during the reaction. The recovered polymer powder had the following properties:

- melt index (2.16 kg load) = 3.74 g/10 mins
- 5 - melt index (5 kg load) = 14.0 g/10 mins
- melt index (21.6 kg load) = 163 g/10 mins
- density = >960 kg/m³
- fines <125µm = 0.83% by weight
- residual iron level = 1.9 ppm

10

Example 2a - Preparation of 2,6-diacetylpyridine bis (2,4,6-trimethylphenyl amine) FeCl₂ complex

The same iron complex was used for Example 2 as for Example 1.

15 Example 2b - Preparation of the catalyst

The equipment used for calcining the silica support was composed essentially of a vertical quartz cylinder, 75cm high and 5cm in diameter, above which was placed a disengagement zone. This calcinator was equipped with a fluidisation grid made from porous quartz and positioned in the lower part of the quartz cylinder. It was also
20 equipped with an electrical resistance heater and a supply of fluidising nitrogen.

Into the calcinator maintained at 60°C and supplied with nitrogen containing less than 2 vpm of water vapour and with a flow rate of 4.7ml/s, were charged 30g of silica sold under the trade name of ES70X by Crosfield Catalysts (Warrington, England). Next the calcinator was heated from 60 to 600°C at a rate of 100°C/h. The silica was
25 then maintained at 600°C for 16 hours in the fluidised state. The silica was then cooled to 25°C and stored under dry nitrogen.

All subsequent operations were carried out in a nitrogen atmosphere. 12.5g of the silica thus dried were suspended in a solution of 0.66g of methylaluminoxane (Witco, Bergkamen, Germany) in 11.1g of toluene. The suspension was agitated for 2 hours at
30 25°C. The silica thus treated was washed five times with 20 ml of toluene. The treated silica was then dried under vacuum.

2.87 g of the treated silica were suspended in 20 ml of toluene and 0.088 g of the above iron complex (see 1a) in 10ml of toluene was added. The suspension was agitated for 17

hours at 25°C. The catalyst thus prepared was washed five times with 20 ml of toluene. The catalyst was then dried under vacuum.

The catalyst thus prepared contained 0.255 % by weight of iron and 2.88 % by weight of aluminium. This corresponds to an atomic ratio Al/Fe of 23.4.

5

Example 2c - Ethylene polymerisation in the gas phase

400g of polyethylene pellets were introduced into a stainless steel reactor of capacity 2.6 litres equipped with a stirrer and maintained under a nitrogen atmosphere. The reactor was heated to 100°C, stirred at 50 rpm and 0.7g of silica previously treated
10 with 1.5mmole/g of triethylaluminium was added as poison scavenger. Next 0.1082g of the above catalyst (see 2b) was added. Then hydrogen was introduced to pressure the reactor to 0.1 MPa. Finally ethylene was fed until a total pressure of 0.8 MPa was obtained. Ethylene was fed to maintain this pressure throughout the reaction. After production corresponding to 50g polyethylene per gram of catalyst, the stirring rate was
15 increased to 100rpm, after the production of 50g of polyethylene the stirring rate was increased further to 150 rpm and after the production of 100g of polyethylene the stirring rate was increased further to 200 rpm.

Polymerisation was allowed to continue for 1 hour and 57 minutes, during which period 224g of polyethylene were produced. This corresponds to an average activity of
20 3424g/mmol Fe/h/bar. The reactor contents were cooled to 25°C and were then recovered from the reactor. A sieve with 2mm diameter mesh was used to separate the polyethylene pellets from the powder formed during the reaction. The recovered polymer powder had the following properties:

- melt index (2.16 kg load) = 0.8 g/10mins
- 25 - melt index (5 kg load) = 3.0 g/10mins
- melt index (21.6 kg load) = 47 g/10mins
- density = >962 kg/m³
- fines <125µm = 0.34% by weight
- residual iron level = 1.2 ppm

30

Example 3 - comparative example

Example 3a - Preparation of 2,6-diacetylpyridine bis (2,4,6-trimethylphenyl amine)

FeCl₂ complex

The same iron complex was used for Example 3 as for Example 1.

Example 3b - Preparation of the catalyst

5 The equipment used for calcining the silica support was composed essentially of a vertical quartz cylinder, 75cm high and 5cm in diameter, above which was placed a disengagement zone. This calcinator was equipped with a fluidisation grid made from porous quartz and positioned in the lower part of the quartz cylinder. It was also equipped with an electrical resistance heater and a supply of fluidising nitrogen.

10 Into the calcinator maintained at 60°C and supplied with nitrogen containing less than 2 vpm of water vapour and with a flow rate of 4.7ml/s, were charged 30g of silica sold under the trade name of ES70X by Crosfield Catalysts (Warrington, England). Next the calcinator was heated from 60 to 600°C at a rate of 100°C/h. The silica was then maintained at 600°C for 16 hours in the fluidised state. The silica was then cooled to 25°C and stored under dry nitrogen.

15 All subsequent operations were carried out in a nitrogen atmosphere. 5.30g of the silica thus dried was suspended in a solution of 1.19g of methylaluminoxane (Witco, Bergkamen, Germany) in 17.07g of toluene. The suspension was agitated for 2 hours at 25°C. The silica thus treated was washed five times with 20 ml of toluene and then dried under vacuum. 1.62g of the treated silica was suspended in 20ml of toluene and 0.045g
20 of the above iron complex (see 1a) in 10 ml of toluene was added. The suspension was agitated for 17 hours at 25°C. The catalyst thus prepared was washed five times with 20ml of toluene. The catalyst was then dried under vacuum.

The catalyst thus prepared contained 0.226% by weight of iron and 4.6% by weight of aluminium. This corresponds to an atomic ratio Al/Fe of 42.3.

25

Example 3c - Ethylene polymerisation in the gas phase

400g of polyethylene pellets were introduced into a stainless steel reactor of capacity 2.6 litres equipped with a stirrer and maintained under a nitrogen atmosphere. The reactor was heated to 100°C, stirred at 50 rpm and 0.7g of silica previously treated
30 with 1.5mmole/g of triethylaluminium was added as poison scavenger. Next 0.131g of the above catalyst (see 3b) was added. Then hydrogen was introduced to pressure the reactor to 0.2 MPa. Finally ethylene was fed until a total pressure of 0.8 MPa was obtained. Ethylene was fed to maintain this pressure throughout the reaction. After

production corresponding to 50g polyethylene per gram of catalyst, the stirring rate was increased to 100rpm, after the production of 50g of polyethylene the stirring rate was increased further to 150 rpm and after the production of 100g of polyethylene the stirring rate was increased further to 200 rpm.

5 Polymerisation was allowed to continue for 4 hours and 28 minutes, during which period 132g of polyethylene were produced. The average activity during this period was 1077g/mmolFe/h/bar. The reactor contents were cooled to 25°C and were then recovered from the reactor. A sieve with 2mm diameter mesh was used to separate the polyethylene pellets from the powder formed during the reaction. The recovered
10 polymer powder had the following properties:

- melt index (2.16 kg load) = 3.17
- melt index (5 kg load) = 12.3
- melt index (8.5 kg load) = 32.5
- residual iron level = 2.2 ppm

15

Example 4 - comparative example

Example 4a - Preparation of 2,6-diacetylpyridine bis (2,4,6-trimethylphenyl amine)

FeCl₂ complex

The same iron complex was used for Example 4 as for Example 1.

20

Example 4b - Preparation of the catalyst

The equipment used for calcining the silica support was composed essentially of a vertical quartz cylinder, 75cm high and 5cm in diameter, above which was placed a disengagement zone. This calcinator was equipped with a fluidisation grid made from
25 porous quartz and positioned in the lower part of the quartz cylinder. It was also equipped with an electrical resistance heater and a supply of fluidising nitrogen.

Into the calcinator maintained at 60°C and supplied with nitrogen containing less than 2 vpm of water vapour and with a flow rate of 4.7ml/s, were charged 30g of silica sold under the trade name of ES70X by Crosfield Catalysts (Warrington, England).
30 Next the calcinator was heated from 60 to 600°C at a rate of 100°C/h. The silica was then maintained at 600°C for 16 hours in the fluidised state. The silica was then cooled to 25°C and stored under dry nitrogen.

All subsequent operations were carried out in a nitrogen atmosphere. 5.14g of

the silica thus dried were suspended in a solution of 0.63g of methylaluminoxane (Witco, Bergkamen, Germany) in 8.99g of toluene. The suspension was agitated for 2 hours at 25°C. The silica thus treated was washed five times with 20 ml of toluene and then dried under vacuum. 2.14g of the treated silica was suspended in 20ml of toluene and 0.074g of the above iron complex (see 1a) in 10 ml of toluene was added. The suspension was agitated for 17 hours at 25°C. The catalyst thus prepared was washed five times with 20ml of toluene. The catalyst was then dried under vacuum.

The catalyst thus prepared contained 0.237% by weight of iron and 5.94% by weight of aluminium. This corresponds to an atomic ratio Al/Fe of 52.0.

10

Example 4c - Ethylene polymerisation in the gas phase

400g of polyethylene pellets were introduced into a stainless steel reactor of capacity 2.6 litres equipped with a stirrer and maintained under a nitrogen atmosphere. The reactor was heated to 100°C, stirred at 50 rpm and 0.7g of silica previously treated with 1.5mmole/g of triethylaluminium was added as poison scavenger. Next 0.1038g of the above catalyst (see 4b) was added. Then hydrogen was introduced to pressure the reactor to 0.2 MPa. Finally ethylene was fed until a total pressure of 0.8 MPa was obtained. Ethylene was fed to maintain this pressure throughout the reaction. After production corresponding to 50g polyethylene per gram of catalyst, the stirring rate was increased to 100rpm, after the production of 50g of polyethylene the stirring rate was increased further to 150 rpm and after the production of 100g of polyethylene the stirring rate was increased further to 200 rpm.

20

Polymerisation was allowed to continue for 3 hours and 58 minutes, during which period 212g of polyethylene were produced. The average activity during this period was 2047g/mmolFe/h/bar. The reactor contents were cooled to 25°C and were then recovered from the reactor. A sieve with 2 mm diameter mesh was used to separate the polyethylene pellets from the powder formed during the reaction. The recovered polymer powder had the following properties:

25

- melt index (2.16 kg load) = 1.29

30

- melt index (5 kg load) = 4.66

- melt index (8.5 kg load) = 11.57

residual iron level = 1.2 ppm

Example 5a - Preparation of 2,6-diacetylpyridine bis (2,4,6-trimethylphenyl amine)FeCl₂ complex

4g of 2,6-diacetylpyridine, 10.32ml of 2,4,6-trimethylaniline and 0.2g of p-toluenesulphonic acid monohydrate were added to 150ml of dry toluene in a 250 ml round bottomed flask. A 20ml Dean-Stark heavy fractionating column and condenser were attached and the mixture heated with stirring to 160°C. The reaction appeared to have gone to completion in about 3 hours. The solution was then evacuated and 100ml of methanol added. 2,6-diacetylpyridine bis (2,4,6-trimethylphenyl amine) precipitated out and was isolated by filtration and washed with 3 aliquots of 10 ml of methanol.

3.19g of iron(II) chloride were weighed into a half litre Schlenk tube and 400 ml of n-butanol added. The suspension was warmed to 90°C and stirred for 3 hours. 10 g of 2,6-diacetylpyridine bis (2,4,6-trimethylphenyl amine) was added as a solid at 80°C. The reaction was stirred for 1 hour then at 25°C over 3 days. The resultant blue suspension was allowed to settle and the supernatant liquid decanted off. 2 washes of the precipitate with 200ml aliquots of toluene were carried out, followed by one wash with 100ml of n-pentane. The residual blue solid was pumped *in vacuo* for 6 hours till dry. It was stored and handled in a nitrogen atmosphere.

Example 5b - Preparation of the catalyst

The equipment used for calcining the silica support essentially comprised a vertical quartz cylinder, 75 cm high and 5 cm in diameter, above which was placed a disengagement zone. This calcinator was equipped with a fluidisation grid made from porous quartz and positioned in the lower part of the quartz cylinder. It was also equipped with an electrical resistance heater and a supply of fluidising nitrogen.

Into the calcinator maintained at 60°C and supplied with nitrogen containing less than 2 vpm of water vapour and with a flow rate of 4.7 ml/s, were charged 30g of silica sold under the trade name of ES70X by Crosfield Catalysts (Warrington, England). Next the calcinator was heated from 60 to 600°C at a rate of 100°C/h. The silica was then maintained at 600°C for 16 hours in the fluidised state. The silica was then cooled to 25°C and stored under dry nitrogen.

All subsequent operations were carried out in a nitrogen atmosphere. 4.06g of the silica thus dried were suspended in a solution of 0.35g of methylaluminoxane (Witco, Bergkamen, Germany) in 17g of toluene. The suspension was agitated for 2 hours at

25°C. The silica thus treated was washed five times with 20ml of toluene. The treated silica was then dried under vacuum.

The silica thus treated was suspended in 20ml of toluene and 0.1492g of the above iron complex (see 5a) was added. The suspension was agitated for 17 hours at 25°C. The catalyst thus prepared was washed five times with 20 ml of toluene. The catalyst was then dried under vacuum.

The catalyst thus prepared contained 0.39 % by weight of iron and 4.01 % by weight of aluminium. This corresponds to an atomic ratio Al/Fe of 21.3.

10 Example 5c - Ethylene polymerisation in the gas phase

400g of polyethylene pellets were introduced into a stainless steel reactor of capacity 2.5 litres equipped with a stirrer and maintained under a nitrogen atmosphere. The reactor was heated to 98°C, stirred at 25 rpm and 0.7g of silica previously treated with 1.5mmole/g of triethylaluminium was added as poison scavenger. Next 0.0825g of the above catalyst (see 5b) was added. Then hydrogen was introduced to pressure the reactor to 0.05MPa. Finally ethylene was fed until a total pressure of 0.8MPa was obtained. Ethylene was fed to maintain this pressure throughout the reaction. After production corresponding to 50g polyethylene per gram of catalyst, the stirring rate was increased to 100rpm, after the production of 50g of polyethylene the stirring rate was increased further to 150 rpm and after the production of 100g of polyethylene the stirring rate was increased further to 200rpm.

Polymerisation was allowed to continue for 1 hour and 15 minutes, during which period 175g of polyethylene were produced. This corresponds to a catalyst yield of 2121g polymer/g catalyst. The average activity during this polymerisation was 2689 g/mM.h.b. The reactor contents were cooled to 25°C and were then recovered from the reactor. A sieve with 2mm diameter mesh was used to separate the polyethylene pellets from the powder formed during the reaction. The recovered polymer powder had the following properties:

- melt index (2.16 kg load) (MI) = 1.50
- melt index (21.6 kg load) (HLMI) = 82.27
- melt index ratio(HLMI/MI) = 54.8
- residual iron level = 2.2 ppm

Example 6a - Preparation of 2,6-diacetylpyridine bis (2,4,6-trimethylphenyl amine)FeCl₂ complex

This was carried out exactly as described in example 5a.

5 Example 6b - Preparation of the catalyst

The equipment used for calcining the silica support was as in Example 5b. Into the calcinator maintained at 60°C and supplied with nitrogen containing less than 2 vpm of water vapour and with a flow rate of 4.7ml/s, were charged 30g of silica sold under the trade name of ES70X by Crosfield Catalysts (Warrington, England). Next the calcinator
10 was heated from 60 to 700°C at a rate of 100°C/h. The silica was then maintained at 700°C for 16 hours in the fluidised state. The silica was then cooled to 25°C and stored under dry nitrogen.

All subsequent operations were carried out in a nitrogen atmosphere. 4.0g of the silica thus dried were suspended in a solution of 0.44g of methylaluminoxane (Witco,
15 Bergkamen, Germany) in 17g of toluene. The suspension was agitated for 2 hours at 25°C. The silica thus treated was washed five times with 20 ml of toluene. The treated silica was then dried under vacuum.

The silica thus treated was suspended in 20ml of toluene and 0.155g of the above iron complex (see 5a) was added. The suspension was agitated for 17 hours at 25°C. The
20 catalyst thus prepared was washed five times with 20ml of toluene. The catalyst was then dried under vacuum.

The catalyst thus prepared contained 0.41 % by weight of iron and 3.61 % by weight of aluminium. This corresponds to an atomic ratio Al/Fe of 18.0

25 Example 6c - Ethylene polymerisation in the gas phase

400g of polyethylene pellets were introduced into a stainless steel reactor of capacity 2.5 litres equipped with a stirrer and maintained under a nitrogen atmosphere. The reactor was heated to 99.5°C, stirred at 25rpm and 0.7g of silica previously treated with 1.5mmole/g of triethylaluminium was added as poison scavenger. Next 0.1184g of
30 the above catalyst (see 6b) was added. Then hydrogen was introduced to pressure the reactor to 0.1MPa. Finally ethylene was fed until a total pressure of 0.8MPa was obtained. Ethylene was fed to maintain this pressure throughout the reaction. After production corresponding to 50g polyethylene per gram of catalyst, the stirring rate was

increased to 100rpm, after the production of 50g of polyethylene the stirring rate was increased further to 150 rpm and after the production of 100g of polyethylene the stirring rate was increased further to 200rpm.

Polymerisation was allowed to continue for 1 hour and 47 minutes, during which
5 period 244g of polyethylene were produced. This corresponds to a catalyst yield of 2061g polymer/g catalyst. The average activity during this polymerisation was 2238g/mM.h.b. The reactor contents were cooled to 25°C and were then recovered from the reactor. A sieve with 2mm diameter mesh was used to separate the polyethylene pellets from the powder formed during the reaction. The recovered
10 polymer powder had the following properties:

- melt index (2.16 kg load) (MI) = 1.77
- melt index (21.6 kg load) (HLMI) = 95.9
- melt index ratio(HLMI/MI) = 54.2
- residual iron level = 2.0 ppm

15

Example 7a - Preparation of 2,6-diacetylpyridine bis (2,4,6-trimethylphenyl amine)FeCl₂ complex

This was carried out exactly as described in example 5a.

20 Example 7b - Preparation of the catalyst

The equipment used for calcining the silica support was as in Example 5b. Into the calcinator maintained at 60°C and supplied with nitrogen containing less than 2 vpm of water vapour and with a flow rate of 4.7ml/s, were charged 30g of silica sold under the trade name of ES70X by Crosfield Catalysts (Warrington, England). Next the
25 calcinator was heated from 60 to 800°C at a rate of 100°C/h. The silica was then maintained at 800°C for 16 hours in the fluidised state. The silica was then cooled to 25°C and stored under dry nitrogen.

All subsequent operations were carried out in a nitrogen atmosphere. 4.0g of the silica thus dried were suspended in a solution of 0.317g of methylaluminoxane (Witco,
30 Bergkamen, Germany) in 17g of toluene. The suspension was agitated for 2 hours at 25°C. The silica thus treated was washed five times with 20ml of toluene. The treated silica was then dried under vacuum.

The silica thus treated was suspended in 20ml of toluene and 0.077g of the above iron

complex (see 5a) was added. The suspension was agitated for 17 hours at 25°C. The catalyst thus prepared was washed five times with 20ml of toluene. The catalyst was then dried under vacuum.

The catalyst thus prepared contained 0.206 % by weight of iron and 2.58 % by weight of aluminium. This corresponds to an atomic ratio Al/Fe of 26.0

Example 7c - Ethylene polymerisation in the gas phase

400g of polyethylene pellets were introduced into a stainless steel reactor of capacity 2.5 litres equipped with a stirrer and maintained under a nitrogen atmosphere. The reactor was heated to 99.8°C, stirred at 25rpm and 0.7g of silica previously treated with 1.5mmole/g of triethylaluminium was added as poison scavenger. Next 0.1150g of the above catalyst (see 7b) was added. Then hydrogen was introduced to pressure the reactor to 0.1MPa. Finally ethylene was fed until a total pressure of 0.8MPa was obtained. Ethylene was fed to maintain this pressure throughout the reaction. After production corresponding to 50g polyethylene per gram of catalyst, the stirring rate was increased to 100rpm, after the production of 50g of polyethylene the stirring rate was increased further to 150 rpm and after the production of 100g of polyethylene the stirring rate was increased further to 200rpm.

Polymerisation was allowed to continue for 1 hour and 23 minutes, during which period 124g of polyethylene were produced. This corresponds to a catalyst yield of 1078g polymer/g catalyst. The average activity during this polymerisation was 3302 g/mM.h.b. The reactor contents were cooled to 25°C and were then recovered from the reactor. A sieve with 2mm diameter mesh was used to separate the polyethylene pellets from the powder formed during the reaction. The recovered polymer powder had the following properties:

- melt index (2.16 kg load) (MI) = 1.42
- melt index (21.6 kg load) (HLMI) = 73.3
- melt index ratio(HLMI/MI) = 51.6
- residual iron level = 1.8 ppm

Example 5a - Preparation of 2,6-diacetylpyridine bis (2,6-dimethyl-4-tertbutylphenyl amine)FeCl₂ complex

4 g of 2,6-diacetylpyridine, 13.5 ml of 2,6-dimethyl-4-tertbutyl-aniline and 0.2 g

of p-toluenesulphonic acid monohydrate were added to 150ml of dry toluene in a 250ml round bottomed flask. A 20ml Dean-Stark heavy fractionating column and condenser were attached and the mixture heated with stirring to 160°C. The reaction appeared to have gone to completion in about 3 hours. The solution was then evacuated and 100ml
5 of methanol added. 2,6-diacetylpyridine bis (2,6-dimethyl-4-tertbutylphenyl amine) precipitated out and was isolated by filtration and washed with 3 aliquots of 10 ml of methanol.

3.19g of iron(II) chloride were weighed into a half litre Schlenk tube and 400 ml of n-butanol added. The suspension was warmed to 90°C and stirred for 3 hours. 12.2g
10 of 2,6-diacetylpyridine bis (2,6-dimethyl-4-tertbutylphenyl amine) was added as a solid at 80°C. The reaction was stirred for 1 hour then at 25°C over 3 days. The resultant blue suspension was allowed to settle and the supernatant liquid decanted off. 2 washes of the precipitate with 200ml aliquots of toluene were carried out, followed by one wash with 100ml of n-pentane. The residual blue solid was pumped in vacuo for 6 hours till
15 dry. It was stored and handled in a nitrogen atmosphere.

Example 8b - Preparation of the catalyst

The equipment used for calcining the silica support was as in Example 5b. Into the calcinator maintained at 60°C and supplied with nitrogen containing less than 2 vpm
20 of water vapour and with a flow rate of 4.7 ml/s, were charged 30g of silica sold under the trade name of ES70X by Crosfield Catalysts (Warrington, England). Next the calcinator was heated from 60 to 600°C at a rate of 100°C/h. The silica was then maintained at 600°C for 16 hours in the fluidised state. The silica was then cooled to 25°C and stored under dry nitrogen.

25 All subsequent operations were carried out in a nitrogen atmosphere. 5.27g of the silica thus dried were suspended in a solution of 0.33g of methylaluminumoxane (Witco, Bergkamen, Germany) in 17.1g of toluene. The suspension was agitated for 2 hours at 25°C. The silica thus treated was washed five times with 20 ml of toluene. The treated silica was then dried under vacuum.

30 1.46 g of the treated silica were suspended in 20 ml of toluene and 0.077g of the above iron complex (see 4a) was added. The suspension was agitated for 17 hours at 25°C. The catalyst thus prepared was washed five times with 20 ml of toluene. The catalyst was then dried under vacuum.

The catalyst thus prepared contained 0.2 % by weight of iron and 3 % by weight of aluminium. This corresponds to an atomic ratio Al/Fe of 31.

Example 8c - Ethylene polymerisation in the gas phase

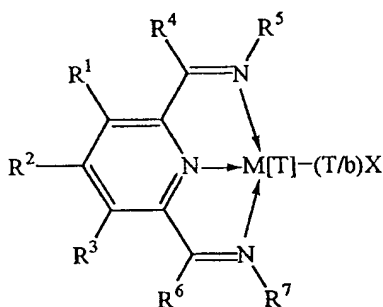
5 400g of polyethylene pellets were introduced into a stainless steel reactor of capacity 2.5 litres equipped with a stirrer and maintained under a nitrogen atmosphere. The reactor was heated to 100°C, stirred at 25 rpm and 0.7g of silica previously treated with 1.5mmole/g of triethylaluminium was added as poison scavenger. Next 0.1214g of the above catalyst (see 8b) was added. Then hydrogen was introduced to pressure the
10 reactor to 0.1MPa. Finally ethylene was fed until a total pressure of 0.8MPa was obtained. Ethylene was fed to maintain this pressure throughout the reaction. After production corresponding to 50g polyethylene per gram of catalyst, the stirring rate was increased to 100rpm, after the production of 50g of polyethylene the stirring rate was increased further to 150 rpm and after the production of 100g of polyethylene the stirring
15 rate was increased further to 200rpm.

Polymerisation was allowed to continue for 1 hour and 23 minutes, during which period 252g of polyethylene were produced. This corresponds to a catalyst yield of 2076g polymer/g catalyst. The reactor contents were cooled to 25°C and were then recovered from the reactor. A sieve with 2mm diameter mesh was used to separate the
20 polyethylene pellets from the powder formed during the reaction. The recovered polymer powder had the following properties:

- melt index (2.16 kg load) (MI) = 2.64
- melt index (5 kg load) = 9.60
- melt index (8.5 kg load) = 23.7
- 25 - melt index (21.6 kg load) (HLMI) = 156
- melt index ratio(HLMI/MI) = 59.1
- residual iron level = 1.2 ppm

Claims

1. Catalyst for the polymerisation of olefins comprising
(1) a complex having the formula (I)



5 Formula (I)

- wherein M is Fe[II], Fe[III], Co[I], Co[II], Co[III], Mn[I], Mn[II], Mn[III], Mn[IV], Ru[II], Ru[III] or Ru[IV]; X represents an atom or group covalently or ionically bonded to the metal M; T is the oxidation state of the metal; b is the valency of the atom or group X; and R¹ to R⁷ are each independently selected from hydrogen, halogen, hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl, substituted heterohydrocarbyl or SiR'₃, where each R' is independently selected from hydrogen, halogen, hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl, substituted heterohydrocarbyl; and when any two or more of R¹ to R⁷ are hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl, said two or more can be linked to form one or more cyclic substituents;
- 10 (2) an activator which is an alkylalumoxane; and
- 15 (3) a support material,
- wherein the atomic ratio of aluminium in the alkylalumoxane (2) to transition metal M in

(1) is from 6:1 to 25:1.

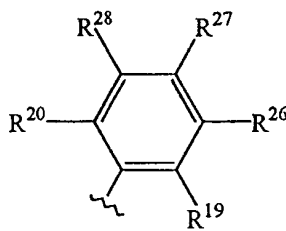
2. Catalyst according to claim 1 wherein the atomic ratio of aluminium to transition metal M is from 8:1 to 22:1.

3. Catalyst according to claim 2 wherein the atomic ratio of aluminium to transition metal M is from 12:1 to 18:1.

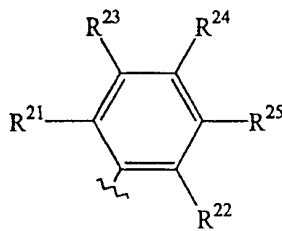
4. Catalyst according to any preceding claim wherein the activator (2) is a (C₁-C₄) alkylalumoxane, and the alkyl group is preferably methyl, ethyl, propyl or isobutyl.

5. Catalyst according to claim 4 wherein the activator (2) is methylalumoxane (MAO) or modified methylalumoxane (MMAO).

6. Catalyst according to any preceding claim wherein R⁵ is represented by the group "P" and R⁷ is represented by the group "Q":



Group P



Group Q

wherein R¹⁹ to R²⁸ are independently selected from hydrogen, halogen, hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl; when any two or more of R¹ to R⁴, R⁶ and R¹⁹ to R²⁸ are hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl, said two or more can be linked to form one or more cyclic substituents.

7. Catalyst according to claim 7 wherein R¹, R², R³, R⁴, R⁶, R¹⁹, R²⁰, R²¹, R²², R²³, R²⁵, R²⁶ and R²⁸ are independently selected from hydrogen and C₁ to C₈ hydrocarbyl, preferably methyl, ethyl, n-propyl, n-butyl, t-butyl, n-hexyl, n-octyl, phenyl and benzyl.

8. Catalyst according to any of claims 1 to 5 wherein R⁵ is a group having the formula -NR²⁹R³⁰ and R⁷ is a group having the formula -NR³¹R³², wherein R²⁹ to R³² are independently selected from hydrogen, halogen, hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl or substituted heterohydrocarbyl; when any two or more of R¹ to R⁴, R⁶ and R²⁹ to R³² are hydrocarbyl, substituted hydrocarbyl,

heterohydrocarbyl or substituted heterohydrocarbyl, said two or more can be linked to form one or more cyclic substituents.

9. Catalyst according to any of claims 6 to 8 wherein R^{24} and R^{27} are each independently fluorine, chlorine, bromine or iodine, or each independently selected from ethyl, n-propyl, iso-propyl, n-butyl, sec-butyl, tert.-butyl, n-pentyl, neopentyl, n-hexyl, 4-methylpentyl, n-octyl, phenyl and benzyl.
10. Catalyst according to any preceding claim wherein the transition metal M is Fe(II), Fe(III) or Co(II).
11. Catalyst according to any preceding claim wherein X is selected from halide, sulphate, nitrate, thiolate, thiocarboxylate, BF_4^- , PF_6^- , hydride, hydrocarbyloxy, carboxylate, hydrocarbyl, substituted hydrocarbyl and heterohydrocarbyl, and β -diketonates.
12. Catalyst according to claim 11 wherein X is selected from chloride, bromide, methyl, ethyl, propyl, butyl, octyl, decyl, phenyl, benzyl, methoxide, ethoxide, isopropoxide, tosylate, triflate, formate, acetate, phenoxide and benzoate.
13. Catalyst according to any preceding claim which comprises 2,6-diacetylpyridinebis(2,6 dimethyl 4, t-butyl anil)FeCl₂, 2,6-diacetylpyridinebis(2,6 dimethyl 4, t-butyl anil)CoCl₂, 2,6-diacetylpyridinebis(2,6 dimethyl 4, t-butyl anil)FeBr₂, 2,6-diacetylpyridinebis(4, t-butyl anil)FeCl₂, and 2,6-diacetylpyridinebis(2,6-dimethyl 4-phenyl anil)FeCl₂ or 2,6-diacetylpyridinebis(2-methyl, 4-fluoroanil) FeCl₂.
14. Catalyst according to any preceding claim which further comprises a neutral Lewis base.
15. Catalyst according to claim 14 wherein the neutral Lewis base is selected from alkenes (other than 1-olefins) or alkynes, primary, secondary and tertiary amines, amides, phosphoramides, phosphines, phosphites, ethers, thioethers, nitriles, esters, ketones, aldehydes, carbon monoxide and carbon dioxide, sulphoxides, sulphones and boroxines.
16. Catalyst according to any preceding claim wherein the support material comprises silica, alumina, MgCl₂ or zirconia, or a polymer or prepolymer comprising polyethylene, polypropylene, polystyrene, or poly(aminostyrene).
17. Process for the polymerisation or copolymerisation of 1-olefins, comprising contacting a monomeric olefin under polymerisation conditions with a catalyst as defined in any preceding claim.
18. Process according to claim 17 comprising the steps of:

a) preparing a prepolymer-based catalyst by contacting one or more 1-olefins with a catalyst, and

b) contacting the prepolymer-based catalyst with one or more 1-olefins,
wherein the catalyst is as defined in any of claims 1 to 16.

- 5 19. Process according to claim 17 or 18 wherein the polymerisation is conducted in the presence of trimethylaluminium (TMA), triethylaluminium (TEA), triisobutylaluminium (TIBA), tri-n-octylaluminium, methylaluminium dichloride, ethylaluminium dichloride, dimethylaluminium chloride, diethylaluminium chloride, ethylaluminiumsesquichloride or methylaluminiumsesquichloride.
- 10 20. Process according to any of claims 17 to 19 wherein the polymerisation is conducted in the presence of hydrogen as a molecular weight modifier.
21. Process according to any one of claims 17 to 20 wherein the polymerisation conditions are solution phase, slurry phase or gas phase.
22. Process according to claim 21 wherein the polymerisation is conducted under gas
15 phase fluidised bed conditions.
23. Process according to claim 21 wherein the polymerisation is conducted in slurry phase in an autoclave or continuous loop reactor.

20

25

30

INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 99/03502

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C08F10/00 C08F4/70

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C08F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 98 27124 A (DU PONT) 25 June 1998 (1998-06-25) cited in the application examples 7,43-46 claims 1-3,45,46 ---	1-23
P,Y	WO 99 12981 A (DORER BIRGIT ANGELIKA ;GIBSON VERNON CHARLES (GB); KIMBERLEY BRIAN) 18 March 1999 (1999-03-18) cited in the application page 44 -page 46; examples 26,27 page 49 -page 59; examples 30-39 claims 1-11,13,14,17-20,23,25,26 --- -/--	1-23

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

3 February 2000

Date of mailing of the international search report

16/02/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Gamb, V

INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 99/03502

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, Y	<p>EP 0 924 223 A (MITSUI CHEMICALS INC)</p> <p>23 June 1999 (1999-06-23)</p> <p>page 124 -page 125</p> <p>page 139, line 12 -page 140, line 17</p> <p>page 150, line 19 -page 152, line 27</p> <p>-----</p>	1-23

INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/GB 99/03502

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9827124 A	25-06-1998	AU 5711098 A EP 0946609 A NO 992930 A US 5955555 A	15-07-1998 06-10-1999 10-08-1999 21-09-1999
WO 9912981 A	18-03-1999	AU 8878398 A AU 2738499 A AU 2738599 A AU 2738699 A AU 2738799 A WO 9946308 A WO 9946302 A WO 9946303 A WO 9946304 A	29-03-1999 27-09-1999 27-09-1999 27-09-1999 27-09-1999 16-09-1999 16-09-1999 16-09-1999 16-09-1999
EP 0924223 A	23-06-1999	NONE	